


Beyond FITT - How Density Can Improve the Understanding of the Dose-Response Relationship Between Physical Activity and Brain Health

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1 **Beyond FITT: How Density Can Improve the Understanding of the Dose-**
2 **Response Relationship Between Physical Activity and Brain Health**

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93

94 **Abstract**

95 Research on physical activity and health, including planned and structured forms such as acute
96 and chronic physical exercise, has focused on understanding potential dose-response
97 relationships. Traditionally, the variables of (i) Frequency, (ii) Intensity, (iii) Time, (iv) and Type
98 (known as the FITT principle) have been used to operationalize the dose of physical activity.
99 In this article, we describe the limitations of FITT and propose that it should be complemented
100 by the underappreciated variable density, which defines the temporal distribution of physical
101 activity stimuli within a single bout of physical activity or between successive bouts of physical
102 activity relative to time spent resting (e.g., in napping/sleeping or sedentary behaviors). Using
103 the field of physical activity and brain health as an example, we discuss challenges and
104 opportunities for further research to use density to improve our understanding of dose-
105 response relationships between physical activity and health-related outcomes.

106

107 **Keywords:** physical exercise, sedentary behavior, brain, cognition, personalized interventions

108

109 **1. Introduction**

110 Physical activity (PA), which includes planned and structured forms such as acute and chronic
111 physical exercise (see Table 1 for definition), is associated with improved brain health across
112 various age groups, and with different health status [1–4]. Regular engagement in PA is
113 beneficial for brain health at multiple levels [5–8], namely (i) the molecular and cellular level
114 (e.g., expression of brain-derived neurotrophic factor [9–15]), (ii) the functional and structural
115 brain level (e.g., brain activity patterns [16–18] or hippocampal volume [19–21]), (iii) the
116 behavioral level (e.g., better cognitive performance [1, 2, 22–30]), and (iv) the risk of adverse
117 health-related events (i.e., lower dementia risk [31–34]). However, the optimal dose of PA,
118 including but not limited to the time point at which PA should be applied or repeated to trigger
119 changes in specific health-related outcomes (i.e., brain health), is not fully understood [6, 8,
120 22, 26, 27, 35, 36].

121 There is currently a need for greater clarity in the definition of the dose of PA (including physical
122 exercise) [37–42]. This extends to the call for a more complete reporting of dose in intervention
123 studies using PA [41, 43–45]. From a practical perspective, elucidating the complex dose-
124 response relationship of PA and health-related outcomes, comprising the interindividual
125 response variability, is an important prerequisite when aiming to maximize the benefits of PA
126 interventions (e.g., on brain health) by individualizing the PA prescription [37, 38, 40, 45–55].

127 Traditionally, the dose of PA has been characterized and prescribed using the FITT principle,
128 an acronym representing: (i) Frequency, (ii) Intensity, (iii) Time (also referred to as duration),
129 and (iv) Type of PA [51, 56–68]. The FITT principle can also be used to retrospectively analyze
130 how the dose of free-living PA (e.g., unplanned and unstructured forms of PA) is associated
131 with health-related outcomes, which can inform recommendations for a specific amount of PA
132 to maintain or improve health. The FITT principle is also commonly used in systematic reviews
133 and meta-analyses when analyzing the dose-response relationship between PA and measures
134 of brain health [26–28, 60]. Some researchers have suggested extending the four elements of
135 the FITT principle by the factors of: (v) Volume (V), which is defined as the total amount of PA
136 spent in a given intensity zone that is typically operationalized as a product of the duration of
137 the acute PA bouts spent in a particular zone of intensity x frequency [57]; and, (vi) Progression
138 (P), which characterizes the gradual and systematic increase of the PA stimulus to maintain
139 overload and, thus, provoke further adaptation(s) [69], into FITT-VP [58, 70]. However,
140 adhering to the FITT-VP principle to prescribe and analyze PA has several disadvantages.

141 First, the FITT-VP principle does not take into account all acute and chronic variables (e.g.,
142 movement frequency) that determine the dose of PA (especially of planned and structured
143 forms such as acute and chronic physical exercise) [37, 38, 40, 71]. Second, the FITT-VP
144 principle does not consider the temporal distribution of PA stimuli within a single bout of PA or

145 between successive bouts of PA relative to the time spent resting, which is conceptualized as
146 density (see definition below) [37, 38, 40]. Third, each component of the FITT-VP principle is
147 treated somewhat independently when in reality variables characterizing PA can be inter-
148 related [37, 71] (e.g., intensity is significantly influenced by other variables such as acute
149 duration [72, 73] and movement frequency [e.g., cadence operationalized as revolutions per
150 minute when using a cycle ergometer] [74, 75]).

151 For example, one study provided evidence that exercise intensity influences the duration
152 individuals can spend in a specific exercise intensity zone [72]. In particular, in healthy younger
153 adults (i) the maximal duration (i.e., defined in minutes) that the participants were able to spend
154 in a given exercise intensity zone during a constant-load exercise test, and (ii) the physiological
155 responses characterizing distinct duration phases during this performance test show a high
156 interindividual variability, while the relative duration (e.g., operationalized as % of maximal
157 duration) was comparable among participants [72]. These findings suggest that a personalized
158 exercise prescription should consider the individualization of the duration spent in specific
159 exercise intensity zones [72, 73].

160 Regarding movement frequency, a study in trained cyclists showed that, at the same exercise
161 intensity, cycling at a higher movement frequency (i.e., 120 revolutions per minute on a cycle
162 ergometer) led to higher physical demands (i.e., operationalized by ratings of perceived
163 exertion, peripheral blood lactate concentration, heart rate, indices of heart rate variability [74],
164 or spectral parameters of the electroencephalography [76]) than cycling at a lower movement
165 frequency (i.e., 60 revolutions per minute) [74, 76]. In addition to the acute differences in
166 physiological markers, there is evidence that in trained cyclists endurance training at different
167 movement frequencies (i.e., high vs. low cadence training for four weeks) may differently
168 influence specific brain measures [77, 78]. In particular, in trained cyclists endurance training
169 at either high or low cadence produces similar improvements in markers of endurance
170 performance (i.e., maximal oxygen uptake and power at the individual anaerobic threshold)
171 [77, 78]. However, training at high cadence led to more pronounced changes in several brain
172 parameters (e.g., reduction in alpha-, beta- and overall-power spectral density [77] or increase
173 in frontal alpha/beta ratio [78] assessed during an incremental exercise test).

174 The above-presented examples highlight the complexity of determining or providing a specific
175 dose of PA and suggest that an oversimplification of dose may hinder accurate prediction and
176 optimization of PA interventions on health [37, 38, 40]. This is also supported by the fact that
177 different PA variables converge in the PA-induced stimulus (i.e., external load) that feeds into
178 the response matrix, where it interacts with non-modifiable factors such as age, sex, or genetic
179 predisposition, and (potentially) modifiable non-PA-related factors such as sleep, nutrition,
180 general stress, and environmental factors, and then triggers specific biological processes that
181 determine the dose (i.e., defined as (a) specific marker(s) of internal load that are involved in

182 biological processes driving the desired changes in outcomes of interest – see Table 1) [37,
 183 38, 40, 71, 79]. Thus accounting for such interrelations of PA variables must not only be
 184 considered when tailoring, programming, or progressing PA interventions [37, 38, 40, 71, 80]
 185 but also as part of the assessment and analytic approaches used.
 186 Consequently, to advance the understanding of the dose-response relationship of PA with
 187 specific domains of health (i.e., brain health [40]), it is necessary to consider additional
 188 variables, such as density, which we will show can allow for a more precise determination of
 189 the dose of PA and provide a more nuanced approach beyond the FITT-VP principle.

190

191 Table 1. Definition of key terms. PA: physical activity; MET: metabolic equivalent of the task;
 192 SB: sedentary behavior

Key terms	
Brain Health	...can be defined as the optimal development and maintenance of brain integrity which encompasses: (i) structural (e.g., hippocampal volume) and functional (e.g., changes in brain activity) brain parameters; (ii) functions that depend on the integrity of the brain, including but not limited to mental health, cognition, and movement; and (iii) the absence of neurological disorders (e.g., dementia). [81, 82]
Dose	...is characterized by three key components: (1) external load (i.e., defined as the work performed by the individual independent of internal characteristics), (2) influencing factors (i.e., all factors [e.g., including environmental factors] that can strengthen or weaken the stimuli of a single bout of PA), and (3) internal load (i.e., defined as the individual and acute physiological, psychological, motor, and biomechanical responses to the external load and the influencing factors during and/or after the cessation of a single bout of PA). Thus, the dose can be operationalized and monitored by using specific indicators of internal load involved in the biological processes that drive the desired changes in outcomes of interest. [37, 40, 79]
Physical Activity (PA)	...can be defined as any muscle-induced bodily movement (e.g., in occupational or leisure time) that results in an increase in the energy expenditure above ~1.5 metabolic equivalents of the task (MET; 1 MET = 1 kcal (4.184 kJ) • kg ⁻¹ • h ⁻¹). This includes planned and structured forms such as acute and chronic physical exercise (see the following definition). PA can be divided into acute (single bout/session of) and chronic (multiple bout/session) PA based on temporal characteristics.” [81, 83–90] Furthermore, PA can be differentiated based on the domains in which it occurs, including recreation/leisure time (such as household), transportation, education, or occupation [87, 88, 91–95].

Physical Exercise	...can be defined as a specific form of PA that is planned, structured, repetitive, and designed to improve or at least maintain the performance in one or more fitness dimensions. Physical exercise can be divided into acute (single bout/session) and chronic (multiple bouts/sessions) based on temporal characteristics, also referred to as physical training [83–86, 88, 89, 91]. In addition, physical exercise is typically performed in recreational/leisure time when it is not part of healthcare service (e.g., rehabilitation) or occupation (e.g., elite athlete). To delimit physical exercise from PA: Physical exercise is always PA, PA is not necessarily physical exercise [96].
Sedentary Behavior (SB)	...can be defined as any waking behavior characterized by a low energy expenditure (≤ 1.5 MET) while sitting or lying down [87–89, 92, 97, 98]. SB is ubiquitous, due to rapid changes in human environmental, economic, social, and technological contexts. Scientifically, SB has been identified as a newer component of the activity spectrum, which can adversely impact health [99–102]. SB can be categorized as cognitively active (e.g., reading) and cognitively passive (e.g., watching television) [81, 103]. For many adolescents and adults, daily time spent sedentary is ≥ 5 hours per day [104–106].

193

194 **2. Method**

195 Given that the German exercise and training variable “Belastungsdichte” [107] (hereafter
 196 referred to as “density”), which has its roots in the field of exercise science, is not well-
 197 recognized internationally, we aimed to improve its accessibility by introducing this variable to
 198 the broader scientific community. In this context, we extend the description and application of
 199 “density” to the field of free-living PA, where it has not previously been applied. As “density” is
 200 underappreciated in the scientific community, we opted to perform a narrative review, since
 201 there is not a large and specific enough literature base to conduct a systematic review (e.g.,
 202 on the role of density of PA on brain health).

203 The author group comprises junior, mid-career, and senior researchers from different
 204 disciplines, and cultural and ethnic backgrounds.

205

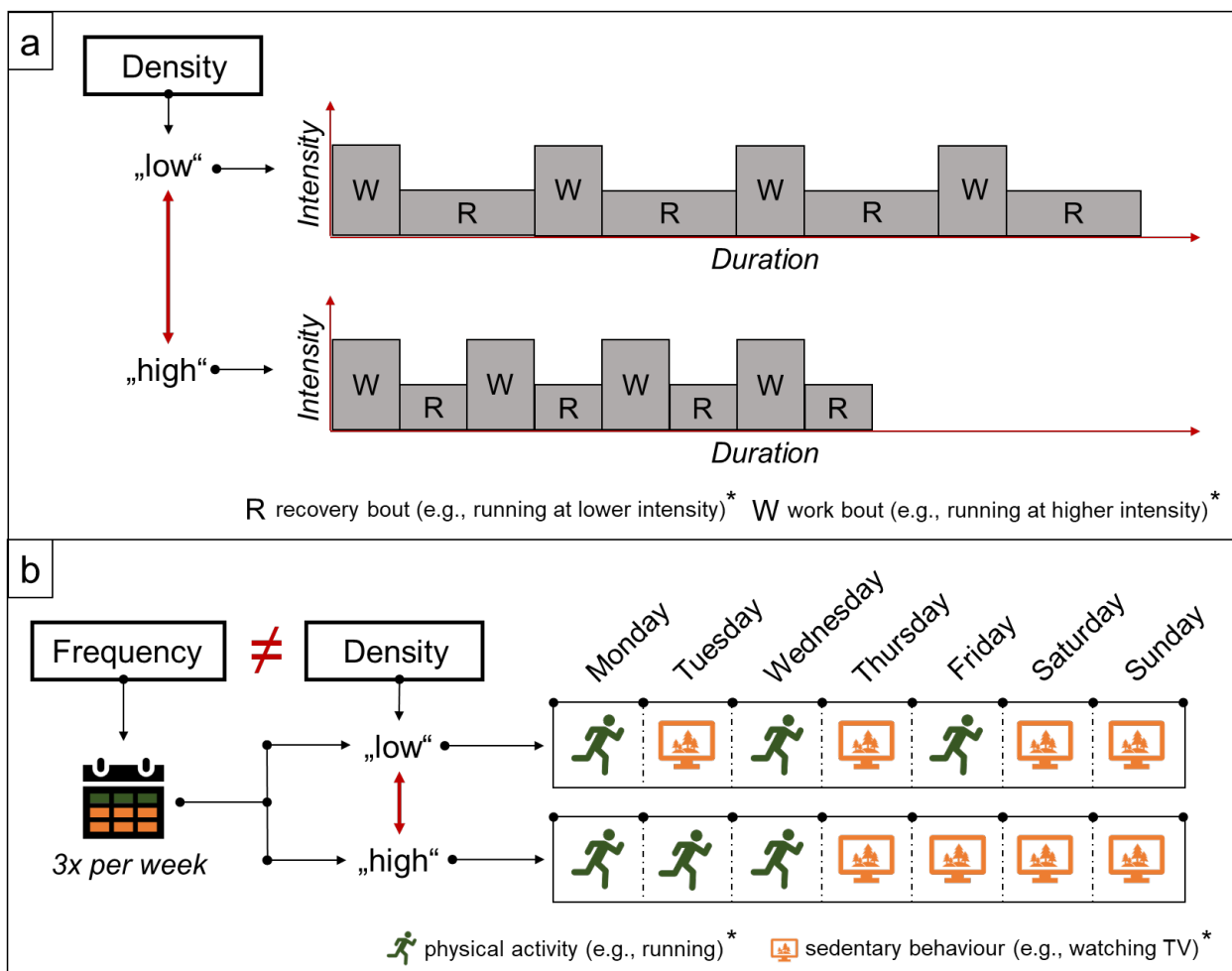
206 **3. Definition of density**

207 Density can be defined as the distribution of PA bout(s) (also referred to as “work bout[s]”) or
 208 portions thereof over a specific time interval (e.g., within a single bout, day, week, month, or
 209 year) in comparison to the time spent resting (also referred to as “rest, recovery or relief bouts”)
 210 [8, 40, 80, 108]. Assuming the characteristics of work bouts remain similar (i.e., are identical
 211 in terms of acute and chronic variables that characterize PA), density is determined by the

212 duration of rest bouts. In other words, density can be modified by changing the duration of
 213 such bouts to adjust the work-rest ratio.

214 In this context, we would like to highlight three important points. First, density is related to the
 215 construct of the work-rest ratio, but differs conceptually in that density is associated with
 216 changing the time spent at rest (i.e., duration of the rest bout[s]), whereas the work-rest ratio
 217 can also be adjusted by increasing the duration of the work bout(s). Second, the variables that
 218 characterize the work bout(s) and the rest bout(s), namely the type of activity, the intensity,
 219 and the duration, need to be considered to gain a more nuanced understanding of the influence
 220 of density and, in turn, the dose-response relationship of PA with measures of brain health.
 221 Third, density needs to be further differentiated based on the temporal context, namely (i) in
 222 acute density (i.e., in the context of acute PA; see Figure 1 a) and (ii) in chronic density (i.e.,
 223 in the context of chronic PA; see Figure 1 b) [37].

224



225
 226 Figure 1: (a) Schematic illustration of different acute densities using an acute bout of physical exercise
 227 in interval mode as an example. In our example, the number of the work bouts (4x) and rest
 228 bouts (4x) is equal whereas the duration of the rest bout in the upper example (i.e., low acute
 229 density; the work-rest ratio of 1:2) is twice as long as in the lower one (high acute density; the
 230 work-rest ratio of 1:1) resulting in a different acute density and, in turn, dose. In this example,
 231 an active rest bout, which is conducted at half of the intensity as the work bout, is selected.

232 The example also illustrates the fact that specific acute variables are interrelated (e.g., acute
233 density, acute duration, and intensity of work and rest intervals). (b) Schematic illustration of
234 the difference between frequency and chronic density in the context of chronic physical
235 activity. The visualization shows that the same frequency (3x physical activity bouts per week)
236 can be distributed differently over a week resulting in a different chronic density and, in turn,
237 dose. The asterisk (*) indicates that other acute (i.e., type of physical activity, intensity, and
238 acute duration) and chronic variables (i.e., chronic duration) that characterize the bout(s) of
239 physical activity are assumed to be constant. Please note that we used sedentary behavior
240 as an example for the rest bout(s). With regard to acute and chronic physical activity, physical
241 activity at a lower intensity than that of the work bout(s), standing, and sleep can be also
242 encompassed by the rest bout(s), depending on the context. Furthermore, the
243 operationalization of chronic density depends on the period of interest (e.g., day, week,
244 month, year).

245

246 **3. Operationalization of acute and chronic density**

247 In the following sections, we propose different approaches to operationalize and analyze
248 density considering the temporal context of PA, the availability and accessibility of population-
249 based datasets, and recent advances in technology to assess PA (i.e., miniaturized wearables
250 to track activities within the 24-hour activity cycle).

251 *3.1 Acute density*

252 As illustrated in Figure 1a, acute density can be operationalized by the duration of the rest
253 bout(s) between the successive work bouts (i.e., in seconds or minutes or relative to the
254 duration of the work bout) within a single session of PA. Thus, a modification of acute density
255 can be achieved by decreasing or increasing the duration of the rest bout(s), resulting in a
256 higher acute work-rest ratio (i.e., higher density) or a lower acute work-rest ratio (i.e., lower
257 density), respectively.

258 *3.2 Chronic density – Simple analysis approaches*

259 The operationalization of chronic density depends on the period of interest (e.g., day, week,
260 month, year). Although chronic density can be operationalized in minutes or hours when
261 several isolated work bouts are performed throughout the day, the operationalization of chronic
262 density is more challenging when longer periods are considered (e.g., week, month, year),
263 especially for unplanned and unstructured forms of PA. To illustrate chronic density in terms
264 of a micro-cycle of one week, consider the following example: if a person is physically active
265 on Monday, Wednesday, and Friday or Monday, Tuesday, and Wednesday, this will result in
266 the same frequency but not the same chronic density within a micro-cycle of one week (see
267 also Figure 1b). More specifically, in the first example shown in Figure 1b, the person is
268 physically active on non-consecutive days (i.e., work bouts spread over a week), whereas in
269 the second example, the person is physically active on consecutive days (i.e., work bouts
270 performed on three consecutive days).

271 Accordingly, a simple approach to studying the influence of different chronic density patterns
272 on brain health is to characterize different groups of individuals based on their chronic density
273 patterns (e.g., a low chronic density group in which individuals performed PA on non-
274 consecutive days versus a high chronic density group in which individuals performed PA on
275 consecutive days – see also Figure 1). For chronic physical exercise, the influence of chronic
276 density on specific measures of brain health can be studied by comparing intervention groups
277 that were instructed to perform physical exercise sessions with different chronic densities (e.g.,
278 a low chronic density group performing physical exercise sessions on non-consecutive days
279 versus a high chronic density group performing physical exercise sessions on consecutive
280 days).

281 *3.3 Chronic density – Sophisticated analysis approaches*

282 Comparable to other studies analyzing the influence of PA patterns (e.g., intensity, and
283 duration of the acute PA bouts) on health-related outcomes (e.g., cognitive performance or
284 cardiometabolic health), the application of more sophisticated approaches using distributional
285 data analysis [109] or machine learning (e.g., via K-means clustering) [110–113] holds some
286 promise for identifying groups of individuals with distinct chronic density patterns. Despite
287 some limitations and challenges (e.g. the need for large sample sizes and, high-dimensional
288 data, the time-consuming nature of training algorithms, and the lack of benchmark data),
289 machine learning-based approaches provide several advantages for the purpose of profiling
290 PA patterns (e.g., more accurate classification and prediction, the possibility of a hypothesis-
291 free/generating approach) [114–118]. Another advantage of machine learning-based
292 approaches is their capacity to handle large, complex, and high-dimensional datasets [114].
293 The ability and flexibility to handle such datasets make machine learning-based approaches
294 well-suited for analyzing the influence of density on specific markers of brain health because
295 density is a more complex variable than other PA variables (e.g. duration). This assumption is
296 supported by the fact that these approaches have already been successfully applied to
297 elucidate the influence of “micropatterns” of PA including intensity and duration (also referred
298 to as bout length) on health-related outcomes such as mortality [119, 120] and cancer
299 incidence [121]. Thus, extending machine learning-based approaches to density is a promising
300 area for future research to elucidate the influence of different chronic density patterns on
301 measures of health in general and brain health in particular.

302 In the context of brain health, the application of such sophisticated classification and analysis
303 techniques may enable the investigation of specific research questions (e.g., *is a low density*
304 *of moderate-intensity PA in older adults more, less, or equally beneficial for brain health than*
305 *having a high density of moderate-intensity PA?*) or to study the association of specific density-
306 related PA patterns, such as the stability of density, with measures of brain health. In this
307 context, we propose that the stability of density is characterized by the periodicity and the

308 fluctuations (variability) that are reflected by the degree of randomness of the duration of the
309 rest bouts between successive work bouts within a given time interval (e.g., day, week, month,
310 year). We suggest that, among other approaches [122], the stability of density can be
311 operationalized by measures used to assess fractal dynamics.

312 Fractal dynamics are characterized by the self-affinity (also referred to as self-similarity or scale
313 invariance) of a given signal (e.g., derived from accelerometers) across time scales [123–127].
314 There is a strong case to be made that fractal dynamics can help to better understand the
315 periodization of chronic physical exercise [128], and several studies have used this approach
316 to analyze physiological data (e.g., frequently applied to heart rate variability data [129–146])
317 or PA patterns [147–150]. In the context of PA, a popular method for assessing fractal
318 dynamics (e.g. of PA [147–150]) is detrended fluctuation analysis (DFA), which is a
319 nonstationary time-series analysis of specific signals (e.g., accelerometer data) that reflects
320 the correlative structure and fractal dimension of signal fluctuations across a range of time
321 scales based on a modified root-mean-square analysis [126, 127, 151–153]. For instance, a
322 study using data from 5097 middle-aged adults showed that greater fractal stability of daily PA
323 (i.e., assessed via a thigh-mounted accelerometer over seven days and reflected in a higher
324 DFA scaling exponent) was associated with better verbal fluency performance in males but not
325 in females [150]. Such sex-specific differences are consistent with the evidence suggesting
326 that sex is an important moderate in the relationship between PA and brain health [47, 48,
327 154–160]. However, whether such findings extend to the chronic density of PA remains a
328 promising area for further investigations.

329 *3.4 Recommendations regarding the assessment of chronic density*

330 To quantify the chronic density of PA, we recommend the application of device-based
331 assessments to complement subjective assessments (i.e., questionnaires) for the following
332 reasons. First, popular questionnaires to assess chronic PA such as the International Physical
333 Activity Questionnaire (IPAQ) only quantify the frequency but not the chronic PA density (i.e.,
334 neither the long form [161] nor the short form [162] of the IPAQ), although some recently
335 developed questionnaires do collect such information (e.g., Daily Activity Behaviours
336 Questionnaire [163–166]). Second, although subjective assessment tools (e.g.,
337 questionnaires) have several advantages (e.g., low burden for participants, cost-effective and
338 convenient administration), they are prone to several sources of bias (e.g., recall bias or social
339 desirability bias) that can confound the estimation of chronic PA patterns [95, 167–169].
340 Device-based assessment tools can circumvent the above-described limitations of subjective
341 assessment tools, but it should be considered that (i) the applied device-based measurement
342 tool needs to be valid and reliable [170–172], and (ii) there is not yet a fully established
343 consensus on the application of device-based measurement tools (e.g., placement and
344 sampling frequency of the device) or on the data processing procedures to obtain specific

345 indices of PA (e.g., minimal length of the epochs, filter, cut-off points, non-wear-time definition)
346 although some recommendations exist [173–175].
347 Furthermore, we recommend combining popular device-based tools such as accelerometers
348 with other sensors (e.g., for environmental light, barometer/altimeter, or geolocation) and
349 digital tools (e.g., smartphones) to allow for the recording of contextual information (e.g.,
350 weather via geolocation at specific time point [176] or type of activity conducted during rest
351 bout(s) via an accelerometer-triggered e-diary [176–182]). The latter approach is also referred
352 to as ambulatory assessment [81, 177, 183, 184]. In addition, regarding the analysis of chronic
353 density in the context of chronic PA, future studies should consider SB and sleep to provide a
354 more holistic understanding of the 24-hour activity cycle on health in general [185–189] and
355 brain health in particular [81, 92, 190–193].

356 *3.5 The potential of density to complement existing analysis approaches of the 24-hour activity* 357 *cycle*

358 Since density specifies the temporal distance between stimuli within or between successive
359 bouts of PA, it can complement other approaches used to analyze the influence of PA patterns
360 within the 24-hour activity cycle on health-related outcomes, namely (i) timing of PA (e.g., time
361 of day on which the PA has been conducted such as in the morning, afternoon or evening
362 [194–196]) and (ii) compositional data analysis (e.g., using the relative time spent in a specific
363 activity [e.g., PA] in relation to the time spent in other activities [e.g., SB or sleep] instead of
364 absolute times spent in a specific activity for analysis [197–204]).

365 In terms of the diurnal impact of PA, PA is an important “Zeitgeber” (time cue) for the human
366 circadian system [205] and thus a critical factor in sleep health, a mediator of the effects of PA
367 on brain health [5, 206]. In this regard, the findings of a recent systematic review suggest that
368 there is currently no consistent evidence in adults as to whether PA conducted at one time of
369 day (e.g., morning) is associated with more pronounced health benefits than PA performed at
370 a different time of day (e.g., afternoon or evening) [194]. In general, PA is associated with
371 better sleep health [207–212], but there is no compelling evidence that PA performed at any
372 particular time of day is superior for promoting sleep health [209, 213, 214] because even
373 acute PA conducted in the evening is not typically detrimental for sleep [215–217] if it is not
374 performed too close before bedtime (≤ 1 hour) [215]. To the best of our knowledge, the timing
375 of PA and its direct relationship with measures of brain health so far has received relatively
376 little attention in empirical studies. The findings from one study suggest that, in adolescents,
377 an acute bout of physical exercise in the morning is more effective in improving behavioral
378 measures of brain health (e.g., global reaction time), compared with the afternoon [218].
379 However, currently (i) there is a lack of studies on the influence of the timing of PA on brain
380 health, and (ii) the evidence on the timing of PA on sleep health, an important mediator of the

381 effects of PA on measures of brain health [5, 206], is less clear. Thus, future research is needed
382 to draw firm conclusions on whether the timing of PA can influence specific measures of brain
383 health differentially [219]. Such future research on the timing of PA is likely to benefit from
384 considering density, which specifies the temporal distance between stimuli within or between
385 successive bouts of PA (e.g., the time between morning and/or evening bouts of PA).
386 Compositional data analysis has been used to investigate the relationship between PA and
387 behavioral measures of brain health in preschoolers [220–222], middle-aged [223], and older
388 adults [224] and has provided valuable insights into the complex relationship between PA and
389 brain health. For example, compared to other activities of the 24-hour activity cycle (e.g., SB
390 and sleep), a loss of time spent in moderate-to-vigorous PA appears to be relatively detrimental
391 to cognitive performance (i.e., cognition composite score) in middle-aged adults, given its
392 smaller relative amount in the 24-hour cycle [223]. Notably, in older adults, longer time spent
393 in light-intensity PA was associated with better inhibitory control (i.e., operationalized by Stroop
394 task performance), especially when accumulated in bouts longer than 10 minutes [224].
395 Comparable to compositional data analysis approaches, a promising area for further
396 investigations is to operationalize density as the relative time spent in work bout(s) (e.g., PA in
397 a specific intensity zone) in relation to the time spent in rest bout(s) (e.g., SB or sleep) to further
398 our understanding of the temporal dynamics of PA and their influence on brain health. Such a
399 better understanding of the temporal dynamics of PA is needed to better inform the
400 individualization of PA interventions [225].

401 *3.6 Interim summary*

402 Taken together, chronic density captures information beyond that provided by frequency,
403 because frequency only specifies the number of PA bouts in a given time interval (e.g., day,
404 week, month, year) but not their distribution within that time interval. Given that the dose of PA,
405 which is influenced by the external load and confounding factors in terms of the acute
406 psychophysiological responses elicited [37, 40], is an important factor in inducing changes in
407 measures of brain health, including cognition [22, 27], it seems reasonable to assume that
408 acute and chronic PA performed at different densities might differentially influence measures
409 of brain health. This latter assumption is also supported by the fact that density is also related
410 to exercise intensity [80, 226, 227] and both acute and chronic density are variables that are
411 important in inducing a specific level of overload and achieving progression [70], both of which
412 are well-known and important factors and principles influencing the dose of PA and therefore
413 the desired outcomes [40, 69]. In the next section, we will discuss the role of density in
414 modifying the dose of PA in more detail.

415

416 **4. Density and the dose of physical activity**

417 Currently, neither the precise dose [6, 8, 22, 26, 27, 35, 36] nor the neurobiological
418 mechanisms that drive the positive effects of acute and chronic PA on brain health are fully
419 understood [5, 6, 23, 40, 228–230]. This knowledge gap extends to the empirical evidence on
420 how density may influence the dose and neurobiological mechanisms that drive brain health.
421 However, our assumption that accounting for density is crucial when aiming to elucidate the
422 dose-response relationship between PA and brain health is supported by evidence from (i)
423 acute PA studies on the temporal dynamics of specific markers of brain health and (ii) studies
424 on glycemic control and brain health in adults with type 2 diabetes, although the latter cannot
425 be readily generalized to healthy adults.

426 *4.1 Temporal dynamics of acute physical activity for brain health*

427 There is some evidence from a meta-analysis that the after-effects of acute physical exercise
428 on cognitive performance are transient, depending on the characteristics of the physical
429 exercises, such as type of physical exercise, intensity, and duration [25]. More specifically,
430 according to this meta-analysis, the greatest effects of acute physical exercises on cognitive
431 performance can be expected 11-20 minutes after the cessation of the acute bout of physical
432 exercises and diminish with longer delays [25]. However, some studies provide evidence that
433 the after-effects of acute physical exercises on specific behavioral measures of brain health
434 (e.g., executive functions) can even persist for up to 30 minutes in healthy younger adults
435 [231–234], 60 minutes in children [235] and younger adults [236], and 90 minutes in healthy
436 younger adults, [237] or even that in healthy younger adults performing acute physical exercise
437 four hours after learning is more beneficial for improving memory performance and
438 hippocampal pattern similarity (i.e., assessed 48 hours later) as compared to performing acute
439 physical exercise immediately after learning the task [238].

440 Based on the paucity of research in this area, the exact time course and moderators (e.g.,
441 acute PA-related factors such as type, intensity, duration, and non-PA-related factors such as
442 age, sex, health status, and fitness level) of the after-effects of acute PA on specific measures
443 of brain health remain somewhat elusive, at least in part due to methodological challenges
444 (e.g., a limited number of follow-up assessments, confounding influence of activities performed
445 between cessation of acute PA and cognitive test administration) [23]. However, based on the
446 above-presented evidence, it is reasonable to assume that considering temporal dynamics of
447 PA - conceptualized as density - has a great potential to add to our understanding of the dose-
448 response relationship of acute PA on specific measures of brain health. More importantly,
449 considering density in future research may help to elucidate the precise time point(s) at which
450 the acute PA stimulus needs to be applied or repeated to prolong the acute PA-related benefits
451 on specific measures of brain health. Such information on the appropriate timing to set a PA

452 stimulus is thus crucial to inform an experimental design and to maximize the effectiveness of
453 PA interventions (e.g., “just-in-time adaptive PA interventions” [239–241]).

454 Several studies support the notion that the density of the PA can be important in optimizing the
455 effectiveness of PA interventions. For example, two studies in healthy younger adults
456 investigated the effects of two repeated acute bouts of high-intensity interval exercise (HIIE,
457 4x 4-minute work bouts at 90% of $VO_{2\text{ peak}}$ interspersed with 3-minute rest bouts at 60% $VO_{2\text{ peak}}$)
458 on inhibitory control (i.e., assessed by the Stroop task every 10 minutes after the cessation
459 of each bout of physical exercise for 5x times) [242, 243]. In both studies, a recovery interval
460 of 60 minutes separated the first bout of acute HIIE from the second bout of HIIE, in which the
461 Stroop task performance was repeatedly assessed [242, 243]. These studies showed that
462 inhibitory control (i.e., reverse Stroop interference score) improved immediately [242, 243] and
463 10 minutes [243] after exercise cessation after the first and second bout acute bouts of HIIE
464 compared to the pretest. However, only after the first acute bout of HIIE the after-effect did
465 persist up to 40 minutes after exercise cessation [242, 243]. In contrast, the executive
466 performance assessed 10 minutes [242] or 20 minutes [243] after the second bout of HIIE was
467 not significantly different from the pretest and was lower than that of the first bout of HIIE when
468 assessments at 20 minutes [242], 30 minutes [242], and 40 minutes [242, 243] (but not 50
469 minutes [242, 243]) after exercise cessation were considered. Collectively, these observations
470 suggest that the acute PA-related effects on inhibitory control were less pronounced in the
471 second bout of HIIE compared to the first bout of HIIE. Hypothetically, such a diminished effect
472 after the second bout of HIIE could be, among other factors, related to the relatively close
473 temporal proximity between the two single bouts of HIIE (i.e., 60 minutes).

474 Based on the observation that the acute PA-induced performance improvements in inhibitory
475 control correlated with changes in blood lactate concentration in both studies [242, 243] and
476 that changes in peripheral blood lactate concentration were significantly lower during and after
477 the second bout of HIIE [243], it seems reasonable to speculate that there is a neurobehavioral
478 relationship between both measures [8, 40, 108, 244, 245]. This assumption is supported by
479 the fact that peripheral blood lactate can cross the blood-brain barrier via monocarboxylate
480 transporters and be utilized as “fuel” for cognitive processes [246–254], which may further
481 explain the positive associations between acute PA-induced blood lactate increases and
482 cognitive enhancement. Indeed, recent studies have reported that changes in peripheral blood
483 lactate concentration are correlated with acute PA-related improvements in cognitive
484 performance [255–257] although it remains somewhat unclear whether blood lactate changes
485 are a mediator of acute PA-induced benefits on cognitive performance because only one study
486 found evidence in favor of this idea [258] while another did not [259].

487 In addition, there is evidence that a change in peripheral blood lactate concentration (e.g.,
488 induced by acute physical exercise [260] or infusion at rest [261]) is associated with a change

489 in the concentration of serum levels of the brain-derived neurotrophic factor (BDNF), an
490 important neurotrophin involved in processes of PA-related neuroplasticity and brain health [7,
491 12, 15, 262–267]. Notably, in younger healthy adults BDNF changes in response to acute PA
492 are correlated with cognitive improvements [268], lending credence to the hypothesis that
493 BDNF is involved in acute PA-induced improvements in behavioral measures of brain health
494 [269]. Such acute PA-triggered effects of BDNF on cognitive performance are likely to be
495 transient, as several studies on the kinetics of BDNF have consistently shown that elevated
496 BDNF levels return to baseline 15-60 minutes after exercise cessation (for review, see [9]),
497 supporting the notion that temporal dynamics (e.g., density) should be considered when
498 examining the effects of acute PA on brain health.

499 Regarding the functional brain level, alterations in cerebral blood flow (CBF) are hypothesized
500 to mediate the acute effects of PA on behavioral measures of brain health [23]. Indeed, some
501 studies provide evidence that acute PA-induced changes in cerebral blood velocity (CBV), a
502 surrogate for CBF that can be operationalized by monitoring middle cerebral artery velocity via
503 transcranial Doppler ultrasound [270–273], correlate with acute PA-induced improvements in
504 behavioral measures of brain health (i.e., executive functioning operationalized by the
505 antisaccade task) [274, 275]. The acute PA-induced increase in CBV can persist for up to 2
506 hours after exercise cessation depending on several factors (e.g., characteristics of the person
507 and the acute bout of PA, methodological factors - for review see [270]) but typically returns to
508 baseline levels relatively shortly after exercise cessation [270, 271] (e.g., 30 minutes - for
509 review see [270]). Comparable to the transient effects of acute PA at the cellular and molecular
510 level (e.g., BDNF), the transient nature of acute PA-related changes at the functional brain
511 level (e.g., CBF) urges future research to consider density as a variable to facilitate our
512 understanding of the neurobiological mechanisms mediating the effects of acute PA on brain
513 health, which is currently relatively scant [5, 23, 229]. Such a better understanding of the
514 temporal dynamics at different levels of analysis [5, 23, 40] (e.g., molecular and cellular levels,
515 such as changes in the noradrenergic and dopaminergic systems [230] or functional levels,
516 such as brain activity or connectivity changes [17, 18]) may yield a more robust understanding
517 of the potential dose-response relationship, which in turn can help to inform future practical
518 applications better.

519 A recent study provided direct evidence that acute density can influence the acute PA-related
520 effects on specific behavioral measures of brain health. In particular, this study used a within-
521 subject crossover design with a pretest-posttest comparison to investigate in healthy younger
522 adults whether the use of different inter-set rest intervals (i.e., 1 minute versus 3 minutes,
523 representing higher and lower acute densities) during an acute bout of low-load resistance
524 exercise (i.e., 40% of a one-repetition maximum, 6x sets of 10x repetitions) can influence acute
525 exercise-induced changes in inhibitory control (i.e., operationalized with the Stroop test) [276].

526 In this study, it was observed that shorter inter-set rest intervals (i.e., 1 minute - high density)
527 improved inhibitory control (i.e., operationalized by a reverse Stroop interference score)
528 immediately, 10 minutes, 20 minutes, and 30 minutes after exercise cessation, whereas such
529 effects were absent for longer inter-set rest intervals (i.e., 3 minutes - lower acute density).
530 Moreover, the improvement in executive functions was greater at 20 and 30 minutes after
531 exercise cessation in the shorter inter-set rest interval condition (i.e., higher acute density)
532 compared with the longer inter-set rest interval condition (i.e., lower acute density) [276]. Thus,
533 the findings of the above-presented study provide strong support for the importance of
534 considering acute density when investigating the dose-response relationship of acute PA with
535 specific measures of brain health.

536 *4.2 Glycemic control and brain health*

537 There is growing evidence that type 2 diabetes, which is characterized by impaired glucose
538 control [277] and poses a public health burden due to its high and still growing worldwide
539 prevalence and related health complications [277–280], is associated with significantly poorer
540 brain health [281–284]. For instance, there is accumulating evidence that type 2 diabetes is
541 associated with reduced structural and functional brain integrity [285–288], lower cognitive
542 performance [285–293], and an increased risk of dementia [294–297]. Given that impaired
543 homeostasis of glucose control is the key feature of type 2 diabetes [277], maintaining “normal”
544 glucose control across the lifespan (e.g., by reducing sedentary behavior and engaging in PA)
545 seems to be an important factor in maintaining brain health, especially in later life stages [298].
546 Indeed, some systematic reviews provide evidence that PA in adults with type 2 diabetes is
547 associated with a positive but weak influence on specific measures of brain health such as
548 cognitive performance, [299–302] although such evidence is not universal, probably due to the
549 heterogeneity of intervention studies in terms of the exercise and training variables
550 characterizing the physical exercise interventions [303].

551 Notably, two small-scaled studies (n = 12 in both studies) in adults with type 2 diabetes showed
552 that interrupting 7 hours of sitting with 3 minutes of light-intensity walking every 15 minutes
553 (i.e., high acute density) was more beneficial for specific measures of glucose control (e.g.,
554 fasting glucose and duration of the dawn phenomenon [304] or post-breakfast and 21-hour
555 glucose control [305]) than interrupting sitting every 30 or 60 minutes (i.e., low acute density)
556 [304, 305]. During the rest periods, the participants had access to a personal computer,
557 internet, and books [304, 305]. Thus, these two small studies in adults with type 2 diabetes
558 provide preliminary evidence that density can influence neurobiological processes (i.e.,
559 glucose control) relevant to brain health [298] which, in turn, supports our idea that considering
560 density is crucial for a more nuanced understanding of the dose-response relationship between
561 PA and measures of brain health. However, the higher density in the above-described studies

562 [304, 305] is also related to a higher frequency of physical exercise bouts, and thus future high-
563 quality studies are needed to (i) disentangle the unique influence of frequency and density on
564 (brain) health-related measures, and (ii) investigate whether different acute and chronic
565 densities of PA might differentially influence specific levels of brain health (e.g., at the
566 molecular and cellular levels such as the release of brain-derived neurotrophic factor).

567 *4.3 Interim summary*

568 Taken together, the evidence on temporal dynamics of specific markers of brain health in
569 response to acute PA and the glucose control - brain health association corroborates our
570 assumption that density is important for advancing our understanding of the dose-response
571 relationship between PA and measures of brain health because it provides crucial information
572 on temporal distribution of PA. More specifically, studying density plays an important role in
573 understanding the minimal and optimal dose by providing information on the minimal and
574 optimal time interval (i.e., rest bout) between PA stimuli within a single bout of PA or successive
575 bouts of PA (i.e., work bouts) being required to maintain or improve specific measures of brain
576 health. Such information on the minimal and optimal time intervals for the delivery of a PA
577 stimulus holds great potential to inform and optimize intervention approaches aimed at
578 promoting PA, such as “just-in-time adaptive PA interventions” [239–241] (e.g., in the context
579 of breaking up prolonged sitting with acute breaks of PA including physical exercise [306–
580 309]).

581

582 **5. Density in relation to other activities of the 24-hour cycle**

583 There is an increasing interest in the scientific community to develop a more holistic
584 understanding of the influence of the 24-hour activity cycle including PA, standing, sedentary
585 behavior (SB), and sleep on health status [185–189] and brain health [81, 92, 190–193].

586 Regarding density, rest bouts are a key construct and may be considered synonymous with,
587 or primary to, time spent in SB when considering waking hours. Epidemiological and
588 experimental evidence shows that sedentary time may influence the relationship between
589 participation in PA and its well-established cardiometabolic health benefits (i.e. highly
590 sedentary individuals may need to do more than the recommended levels of PA to offset the
591 detrimental effects of sedentary behavior) [99–101, 310]. Experimental evidence provides
592 compelling insights into the potential for “exercise resistance” [100]. Coyle and colleagues
593 showed that when acute physical exercise was preceded by a prolonged period of SB,
594 postprandial metabolic responses and metabolic benefits were significantly attenuated [311–
595 313]. More specifically for brain health, the effect of physical exercise on cognitive function is
596 altered by subsequent exposure to prolonged sitting versus breaks in sitting [306], and

597 emerging evidence shows that different types of SB, namely passive and mentally active SB,
598 could be differentially associated with brain health [81, 103, 314]. For instance, previous
599 studies have indicated that mentally active SB (e.g., reading or using a computer) can benefit
600 measures of brain health (for review see [81, 103, 314]). A growing body of evidence suggests
601 that the consequences of too much time spent in SB are distinct from those of too little PA with
602 respect to cardiometabolic health [100] and brain health [81, 101, 105]. This reinforces the
603 utility of considering SB as a mechanism for the importance of density as a key new element
604 to complement the FITT-VP principle.

605 Given that the duration and the characteristics of the rest bout(s) are the key elements in
606 defining density, considering sleep is important in understanding how the temporal distance
607 between successive bouts of PA can influence measures of brain health, especially when
608 tracking and analyzing free-living PA over longer periods (e.g., a week, month, or year). There
609 is growing evidence that sleep (i.e., often operationalized as time in bed) can mediate and/ or
610 moderate the effect of PA on brain health [193, 219, 315, 316]. For example, several cross-sectional
611 studies provide evidence that (i) older adults with poor sleep efficiency (i.e., percent of the time
612 in bed spent asleep) benefit most from PA in terms of global cognition [317], (ii) sleep efficiency
613 mediates the relationship between PA and working memory, task switching, verbal ability and
614 fluency, and memory recall in a mixed sample of younger and older adults [21], (iii) better
615 subjective sleep quality mediates the relationship between PA and verbal fluency, immediate
616 recall, and delayed recall [318] or working memory [319] in middle-aged and older adults, and
617 (iv) subjective sleep quality and sleep efficiency mediate the relationship between PA level and
618 inhibitory control in younger adults [320]. A 6-month intervention study, in which cognitively
619 healthy older adults performed moderate- or high-intensity interval exercise twice a week,
620 reported that participants in the moderate-intensity group, who had poorer sleep efficiency at
621 baseline, showed greater exercise-induced improvements in episodic memory and global
622 cognition [321].

623 Collectively, the above-presented evidence supports the idea that consideration of all activities
624 in the 24-hour activity cycle [81, 92, 190–193, 316] is necessary to improve our understanding
625 of the influence of specific lifestyle-related factors on brain health. This assumption is
626 reinforced by emerging evidence suggesting that (i) other activities of the 24-hour cycle that
627 can contribute to or constitute the rest bout(s), such as free-living standing activity [322] and
628 light-intensity PA [29], are positively associated with behavioral measures of brain health, and
629 (ii) activities such as SB and sleep, which are typical activities of a rest bout(s), interact with
630 each other with respect to brain health, as an observational study showed that sleep problems
631 mediated the detrimental associations of passive SB with depression [323]. To this end,
632 complementing the 24-hour activity cycle approach with density may enable even more

633 nuanced insights into its health effects by improving the characterization and thus our
634 understanding of the dose of PA.

635

636 **6. The current state of evidence and future directions**

637 The role of density as an important variable can be considered helpful when investigating dose-
638 response relationships of PA with key health-related outcomes (e.g., brain health). For brain
639 health, the current evidence indicates that (i) acute density is typically not considered when
640 analyzing the influence of acute bouts of PA on cognitive performance (e.g., as a moderator
641 variable) [23, 25, 324–326], (ii) chronic density is often not reported in studies investigating the
642 influence of chronic PA on brain health [8, 327], (iii) chronic density is absent in moderator
643 analyses in recent systematic reviews and meta-analyses investigating the influence of chronic
644 PA on cognitive performance [22, 26], and (iv) chronic density is typically not mentioned in
645 recommendations (e.g. from the World Health Organization) and policies aimed at reducing
646 the risk of cognitive decline and dementia by lifestyle changes (e.g., via PA) [328]. Such an
647 absence of density in the literature, analyses of the dose-response-relationships, and
648 recommendations of official bodies could lead to the assumption that (i) acute and chronic
649 density are unimportant variables or (ii) that researchers studying the effects of PA on
650 measures of brain health are unaware of the importance of density.

651 Given that other fields of research have begun to recognize the influence of the distribution of
652 PA across a week (e.g., the “weekend warrior” pattern characterized by $\leq 2x$ bouts [329–336]
653 or 1x bout [337] of PA per week) and the interrelated impacts of PA, sleep, and SB [81, 92,
654 100, 101, 186–189, 191–193, 316], density is an excellent candidate determinant of brain
655 health effects that should not be overlooked when analyzing the dose-response relationship
656 within the context of PA-related benefits on measures of brain health. To simulate future
657 research, we highlight in the following two sections further directions for observational and
658 intervention studies on the influence of PA density on measures of brain health.

659 *6.1 Observational studies*

660 Other research fields have started to analyze observational and population-based data in
661 adults regarding the influence of achieving the amount of PA recommended by the World
662 Health Organization (i.e., ≥ 150 minutes of moderate- or ≥ 75 minutes of vigorous-intensity PA
663 per week [88, 89]) in $\leq 2x$ bouts per week (i.e., denoted as “weekend warrior”) or $\geq 3x$ bouts per
664 week on health-related outcomes such as the risk of mortality [329–331], risk of cardiovascular
665 events [336], prevalence and health aspects associated with the metabolic syndrome (e.g.,
666 adiposity, hypertension) [332, 334], or risk of mental disorders [333]. Although none of the
667 above-mentioned studies considered chronic density, because they did not account for the

668 temporal distance between the successive bouts of PA into account, all provided evidence that
669 achieving the recommended amount of PA in $\leq 2x$ bouts per week has a comparable influence
670 on health-related outcomes as achieving this amount in $\geq 3x$ bouts per week [329–334, 336].

671 Whether such observation extends to measures of brain health, given the moderating role of
672 the acute and chronic density of PA is considered, is a promising area for further investigations.
673 In this regard, we would like to acknowledge that all activities of the 24-hour activity cycle (i.e.,
674 PA, sedentary behavior, and sleep) should be considered for a more nuanced understanding
675 of the dose-response relationship between PA and health in general [185, 186] and brain
676 health in particular [81, 92, 191, 192]. In the context of acute and chronic density, we reiterate
677 that the characteristics that define the work bout(s) and rest bout(s) must be considered when
678 analyzing density (i.e., type of activity, intensity, and duration). This assumption is supported
679 by emerging evidence showing that the characteristics of activities that are primarily involved
680 in the rest bout(s) can influence brain health differentially. More specifically, there is evidence
681 that the type of SB can moderate the effects of SB on brain health because cognitively active
682 SB (e.g., reading) is positively associated with brain health, whereas cognitively passive SB
683 (e.g., watching TV) did not confer such benefits [81, 103, 314, 338].

684 In addition, from a public health perspective, a key distinction is made between active and
685 passive (sedentary) occupations [339]. In this context, analyzing the influence of acute and
686 chronic density on measures of brain health might be especially relevant for health-related
687 research in individuals with professions that require performing substantial occupational PA at
688 higher intensities in relatively short time intervals (e.g., construction workers, or farmers)
689 versus desk-based workers. Considering density in addition to traditional exercise variables
690 (e.g., FITT-VP principle) may enhance our understanding of the “physical activity paradox”
691 (i.e., occupational PA has less clear or no health benefits compared to leisure-time PA) [340–
692 344] and the identification of “sweet spots” (e.g., individualizing leisure time PA
693 recommendations by considering occupational PA levels) [187] which in turn can help to better
694 inform future public health interventions. The latter assumption is reinforced by the fact that
695 individuals with a lower socioeconomic position (i.e., lower educational qualifications,
696 occupational class, income, or living in a deprived area), as compared to those with a higher
697 socioeconomic position, showed different characteristics concerning their 24-hour activity
698 cycle since they spent more time standing, moving, and walking but less time sitting during
699 weekdays while on weekends these patterns were reversed [345]. Notably, those with higher
700 socioeconomic positions engaged in higher levels of physical exercise-like activities (i.e.,
701 running, cycling, and inclined walking) and less time lying regardless of the day of the week.
702 These findings suggest that socioeconomic disadvantages are mirrored in 24-hour activity
703 cycle patterns [345]. Such an observation is of particular relevance for future studies on PA
704 and brain health given that in adults a lower socioeconomic position is negatively associated

705 with different markers of brain health (e.g., lower cognitive function and higher cognitive decline
706 [346–356], higher dementia risk [355, 357–360], less favorable brain structure outcomes [353,
707 354, 360, 361]). Future well-designed research is needed for more robust conclusions in this
708 direction [362, 363] and may benefit from considering the 24-hour activity cycle [191] including
709 the density of PA.

710 *6.2 Intervention studies*

711 In addition to the examination of density in observational studies, we also recommend that
712 acute and chronic density should be considered in the prescription of PA intervention studies
713 to improve the standardization of reporting, the determination of the dose, and the
714 comparability across studies. Although there is evidence that a higher frequency (i.e., 5-7 PA
715 sessions per week), which is probably also mirrored in a higher chronic density, is more
716 beneficial for improving cognitive performance in adults older than 50 years (i.e., double the
717 effect size; 0.69 vs 0.32) than a lower frequency (i.e., 1-2 PA sessions per week) [27], providing
718 information on acute and chronic density can be especially relevant for interventions with lower
719 levels of direct supervision (e.g., home- and technology-based interventions using
720 exergames). For example, in home-based studies using exergames and providing only general
721 supervision, partial direct supervision, or even no supervision (for more information on
722 supervision please see [364, 365]), older adults are typically instructed to achieve a certain
723 duration of physical exercise over a week but are often allowed to self-select the frequency of
724 the acute PA bouts [366–373]. Such studies have documented that older participants who are
725 highly motivated can exceed the recommended training frequency and/or perform multiple
726 acute PA bouts throughout the day [368, 373–375]. This may result in insufficient rest time,
727 which is perhaps less than optimal for the materialization of adaptation processes (i.e.,
728 consolidation). The above theoretical assumption is supported by (i) an experimental study
729 showing that in younger adults too much consecutive computer-based training can be
730 detrimental to learning performance (i.e., accuracy of motion discrimination) [376] and (ii) a
731 systematic review observing that cognitive performance declines when endurance athletes are
732 overreached or overtrained [377]. These latter findings support the assumption that acute and
733 chronic density should be considered when prescribing and monitoring physical exercise
734 interventions aimed at promoting brain health.

735 In particular, acute and chronic density are important variables in the organization of physical
736 exercise, namely the periodization and programming of physical exercise sessions, because
737 they characterize the dose by defining the duration of rest bout(s) within a single bout of
738 physical exercise or between successive bouts of physical exercise (i.e., work bouts). Whereas
739 periodization is the temporal organization (i.e., macro-management) of the characteristics of
740 physical exercise sessions (e.g., purposeful adjustment of variables such as exercise intensity
741 and volume for progression) and application of training principles [37, 40, 128, 378–380],

742 programming is defined as the micro-management of physical exercise that includes, but is
743 not limited to, the organization of exercise and training variables (e.g., type of physical
744 exercise, exercise intensity, exercise duration, and acute and chronic density) [40, 378, 380].
745 Thus, acute density is especially relevant for programming acute physical exercise sessions
746 in which the physical exercises are performed in interval mode or a set structure because acute
747 density defines the rest duration between the work bouts (e.g., also referred to as intervals or
748 repetitions), between interval series or sets, or between different physical exercises [80, 227,
749 379]. As shown in Figure 1, acute density can be manipulated to alter the acute PA stimulus
750 by decreasing or increasing the duration of rest between successive work bouts.

751 From the perspective of PA promotion, density can also complement newer approaches to
752 foster PA, such as “vigorous intermittent lifestyle physical activity” (VILPA) [381, 382] and
753 “exercise snacks” [381–384]. While VILPA has been empirically defined as vigorous bouts of
754 incidental PA lasting up to 1 or 2 minutes [119, 121], the term “exercise snacks” has been more
755 loosely defined as single planned bouts of physical exercise that typically (i) lasts ≤ 1 minute,
756 (ii) occur multiple times throughout the day, and (iii) are performed at a vigorous intensity [382–
757 384]. Regarding the VILPA and “exercise snacks” concepts, the variable density as a
758 characteristic defining the dose can help to more precisely elucidate the influence of different
759 rest durations between the short work bouts (e.g., performed at the vigorous intensity and
760 conceptualized in the VILPA and “exercise snacks” approach or at other intensities in the
761 context of free-living PA such as light- or moderate-intensity PA) on health-related parameters
762 (e.g., brain health). However, it is worth noting that for a purposeful modification of density, the
763 interrelation with other exercise variables needs to be considered (e.g., implementation of
764 passive or active rest periods, exercise intensity, and duration of work and rest bouts) [37, 38,
765 40, 71, 80].

766

767 **7. Limitations**

768 In this article, we advocate the extension of the FITT-VP principle from a physiological
769 perspective by proposing density as an additional variable that allows for a more fine-grained
770 characterization of the dose of PA. However, the following limitations need to be
771 acknowledged. First, it should be noted that others have already advocated for complementing
772 FITT from a psychological perspective by integrating an additional “F” representing “fun” as an
773 umbrella term for psychological factors such as affective valence and enjoyment of PA [385]
774 to reflect that these factors are important determinants of engagement and adherence to PA
775 [386–391]. Second, although we provide in this article a strong theoretical rationale that
776 complementing FITT-VP by the variable density will improve our understanding of the dose-
777 response relationship between PA and health-related outcomes, we wish to emphasize that

778 the precise characterization or prescription of a specific PA dose will remain a considerable
779 challenge because of the myriad of (i) non-modifiable factors (e.g., age, sex, genetics), (ii)
780 potentially modifiable non-PA-related factors (e.g., diet, sleep, stress, environmental
781 conditions), and (iii) modifiable PA-related factors (e.g., type of PA, intensity, duration,
782 movement frequency), which include but are not limited to setting (e.g., home-based or center-
783 based, and indoor or outdoor), method of delivery (e.g., in-person or online), level of
784 supervision (e.g., no supervision, general supervision, direct supervision) and social interaction
785 (e.g., individual or group-based), that can influence the dose and individual
786 psychophysiological response(s) to PA [37, 38, 40, 45, 54, 71, 364]. In other words, adding
787 density to FITT-VP is another piece of the puzzle to better characterize the dose of PA and, in
788 turn, disentangle its influence on specific health-related outcomes.

789

790 **8. Conclusions**

791 In summary, we have provided an overview of the implications and the potential of addressing
792 the density of PA as a variable that has been under-recognized when studying the relationship
793 between PA and health-related outcomes, using the field of brain health as an example. In
794 view of an increasing interest in understanding the dose of PA including but not limited to
795 “micropatterns” assessed using high-resolution wearable data [119, 120, 392], density is a
796 variable that can complement the traditional concept (i.e., the FITT-VP principle) by
797 considering an additional element - the temporal distribution of PA stimuli within a single bout
798 of PA or between successive bouts of PA relative to the time spent resting. We propose a
799 definition for density and approaches for operationalizing it which, in turn, may allow for a more
800 precise determination of the dose of PA for improved health effects and the prevention and
801 treatment of chronic disease. Considering that an explicit focus on the density variable has
802 been largely absent from research to date, investing greater effort in understanding it will add
803 fruitful nuance to identifying the dose-response relationship between PA and health-related
804 outcomes (e.g., brain health), and thus has the potential to provide important information on
805 the optimal and minimal beneficial doses of PA.

806 **Declarations**

807

808 **Authors' Contributions**

809 F.H.: conceptualization, writing – original draft, visualization; L.Z., P.T., P.M., R.F., Q.Y., T.L.-
810 A., C.H., A.F.K., K.E., B.C., Y.C., M.H., Z.Z., T.I., K.K., S.A., Y.G., J.C., M.H., M.H., Z.C., D.M.,
811 V.F., D.R., E.S., M.W. N.O., S.L., H.B.: writing – review & editing. T.G.: writing – review &
812 editing, supervision. All authors read and approved the final version of the manuscript.

813

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816

817 **Ethics approval and consent to participate**

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819

820 **Consent for publication**

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825

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827 The authors declare no conflict of interest or competing interests.

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831

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